

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
ORR-3	15	Estimate responses 75	0.25	Estimated 281.25
ORR-4	15	Estimate responses 119	1.25	Estimated 2231.25

Estimated Total Annual Burden Hours: 2512.5.

In compliance with the requirements of Section 506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. Email address: infocollection@acf.hhs.gov. All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2012-19418 Filed 8-7-12; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Proposed Collection; Comment Request; Semi-Annual and Final Reporting Requirements for Discretionary Grant Programs

AGENCY: Administration for Community Living, HHS.

ACTION: Notice.

SUMMARY: The Administration for Community Living (ACL) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection requirements relating to the continuation of an existing collection for Performance Progress Reports previously approved for discretionary grants funded by the U.S. Administration on Aging (AoA), which is now a part of ACL.

DATES: Submit written or electronic comments on the collection of information by October 9, 2012.

ADDRESSES: Submit electronic comments on the collection of information to: lori.stalbaum@aoa.hhs.gov. Submit written comments on the collection of information to Lori Stalbaum, Administration on Aging, Washington, DC 20201 or by fax to Lori Stalbaum at 202-357-3469.

FOR FURTHER INFORMATION CONTACT: Lori Stalbaum at 202-357-3452 or lori.stalbaum@aoa.hhs.gov.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501-3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. "Collection of information" is defined

in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency request or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, ACL is publishing notice of the proposed collection of information set forth in this document. With respect to the following collection of information, ACL invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of ACL's functions, including whether the information will have practical utility; (2) the accuracy of ACL's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques when appropriate, and other forms of information technology.

The Administration for Community Living (ACL) plans to continue an existing approved collection of information for semi-annual and final reports pursuant to the requirements of its discretionary grant programs. Through its discretionary grant programs, ACL supports projects for the purpose of developing and testing new knowledge and program innovations with the potential for contributing to the independence, well-being, and health of older adults, people with disabilities across the lifespan, and their families and caregivers. Deliverables required by ACL of all Title IV grantees are semi-annual and final reports, as provided for in the Department of Health and Human Services regulations, 45CFR Part 74, Section 74.51. These Title IV grantee performance reporting requirements can be found on ACL's Web site at <http://>

www.aoa.gov/AoARoot/Grants/Reporting_Requirements/docs/FinalReportHandbook.doc. ACL estimates the burden of this collection of information as follows: *Frequency:* Semi-annually with the Final report taking the place of the semi-annual report at the end of the final year of the grant. *Respondents:* States, public agencies, private nonprofit agencies, institutions of higher education, and organizations including tribal organizations. *Estimated Number of Responses:* 600. *Total Estimated Burden Hours:* 12,000.

Dated: August 3, 2012.

Kathy Greenlee,
Administrator and Assistant Secretary for Aging.

[FR Doc. 2012-19453 Filed 8-7-12; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2011-N-0608]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; MedWatch: The Food and Drug Administration Medical Products Reporting Program

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled "MedWatch: The Food and Drug Administration Medical Products Reporting Program" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50-400B, Rockville, MD 20850, 301-796-3794, Jonnalynn.Capezzuto@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: On June 28, 2012, the Agency submitted a proposed collection of information entitled "MedWatch: The Food and Drug Administration Medical Products Reporting Program" to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the

information collection and has assigned OMB control number 0910-0291. The approval expires on June 30, 2015. A copy of the supporting statement for this information collection is available on the Internet at <http://www.reginfo.gov/public/do/PRAMain>.

Dated: August 3, 2012.

David Dorsey,
Acting Associate Commissioner for Policy and Planning.

[FR Doc. 2012-19377 Filed 8-7-12; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2012-N-0793]

Request for Nominations of Specific Drug/Biologic Product(s) That Could Be Brought Before the Food and Drug Administration's Pediatric Subcommittee of the Oncologic Drugs Advisory Committee

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; request for product nominations.

SUMMARY: The Food and Drug Administration's (FDA) Office of Hematology and Oncology Products invites the public to suggest one or more specific drug or biologic products that could be brought before the December 4, 2012, Pediatric Subcommittee of the Oncologic Drugs Advisory Committee (ODAC). The number of drugs studied for use in pediatric patients is growing, and we see a reduction in off-label use. However, we would like to improve current and future pediatric product development by focusing on products whose development would benefit the most from the attention of an advisory committee. The company developing a product that is brought before the committee will be given the unique opportunity to present proposed pediatric studies in the United States, share their plans for global pediatric development, and hear discussions by the Pediatric Subcommittee on possible directions for their current or future pediatric oncology product development.

DATES: Nominations must be received by September 4, 2012, to receive consideration for inclusion. Nominations received after this date will receive consideration for future meetings of the Pediatric Subcommittee of the ODAC.

ADDRESSES: Email nominations to Christine.Lincoln@fda.hhs.gov, and please include the subject line "Suggested Product for 2012 Pediatric Oncology Subcommittee of ODAC."

FOR FURTHER INFORMATION CONTACT: Christine Lincoln, RN, MS, MBA, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 22, Rm. 2206, Silver Spring, MD 20993, 301-796-4117, Christine.Lincoln@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: The Food and Drug Administration (FDA) Advisory Committees are an important, transparent interface that allows the Agency to include the public in its decision-making processes. Significant public health and safety issues are brought before these committees for deliberation, and the meetings bring together both experts with state-of-the-art knowledge and members of the public with relevant personal experiences. This broad participation gives FDA a unique perspective as it seeks to assure the safety, efficacy, and security of FDA-regulated products.

Additional information about the prior November 2, 2011, Pediatric Oncology Subcommittee of the Oncologic Drugs Advisory Committee may be found on FDA's Web site at: <http://www.fda.gov/AdvisoryCommittees/Calendar/ucm274396.htm>.

Dated: August 2, 2012.

Leslie Kux,
Assistant Commissioner for Policy.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Statement of Organization, Functions and Delegations of Authority

This notice amends Part R of the Statement of Organization, Functions and Delegations of Authority of the Department of Health and Human Services (HHS), Health Resources and Services Administration (HRSA) (60 FR 56605, as amended November 6, 1995; as last amended at 77 FR 46098-46099 dated August 2, 2012).

This notice reflects organizational changes in the Health Resources and Services Administration (HRSA). Specifically, this notice updates the functional statement for both the Office of Operations (RB) and the Office of Management (RB4) to include the human resources function for HRSA;